Erdheim-Chester disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Erdheim-Chester disease. ORPHA:35687

Erdheim-Chester disease (ECD), a non-Langerhans form of histiocytosis, is a multisystemic disease characterized by various manifestations such as skeletal involvement with bone pain, exophthalmos, diabetes insipidus, renal impairment and central nervous system (CNS) and/or cardiovascular involvement.